

Taxonomic revision of *Sinoeugnathus kueichowensis* (Halecomorphi, Holostei) from the Middle Triassic of Guizhou and Yunnan, China

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Abstract The previously alleged ‘eugnathid amiiform’ *Sinoeugnathus kueichowensis* is a small-sized halecomorph from the Middle Triassic (Ladinian) marine deposits of Guizhou and Yunnan, China. A morphological redescription and taxonomic revision of this taxon are provided based on a detailed examination of 15 new specimens. Among them, IVPP V24315 (standard length = 64 mm) is appointed as the neotype, given that the holotype is missing. Studies of these specimens revealed some morphological details previously undescribed or misidentified for this taxon, including a hatchet-shaped antorbital, two broad suborbitals, a sensory canal in the maxilla, and three pairs of extrascapulars. For the first time, *Sinoeugnathus* was incorporated into an analysis of halecomorph phylogeny, and the results recover it as the sister taxon of the Anisian *Subortichthys* from Luoping, Yunnan, and both are grouped with two Ladinian genera *Allolepidotus* and *Eoeugnathus* from the Monte San Giorgio area into a monophyletic group (namely Subortichthyidae fam. nov. herein) at the base of Ionoscopiformes. This taxonomic reassessment of *Subortichthys* provides new insights into the phylogeny and paleogeographic evolution of Ionoscopiformes.

Key words Guizhou and Yunnan, Middle Triassic, Ionoscopiformes, Halecomorphi, osteology, phylogeny

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1 Introduction

Holostei includes two divisions, Halecomorphi (e.g., bowfin) and Ginglymodi (e.g., gars) (Cavin, 2010; Grande, 2010; Xu, 2019). Recent morphological and molecular studies consistently recover the Holostei as a monophyletic group sister to Teleostei (e.g., carps and salmons), but the origin and early evolution of this clade remain obscure because of insufficient

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studies of fossil records (Hurley et al., 2007; Near et al., 2012; López-Arbarello and Sferco, 2018). Paleozoic holosteans are scarce and taxonomically ambiguous. The Late Permian (Wuchiapingian) *Acentrophorus* is likely the earliest holostean but still needs a restudy and a formal analysis (Gill, 1923; Friedman, 2015). The earliest, unambiguous holosteans are represented by the Early Triassic parasemionotiform halecomorphs (Olsen, 1984; Grande and Bemis, 1998). In the Middle Triassic, holosteans underwent a rapid radiation, represented mainly by ionoscopiform halecomorphs (Xu et al., 2014; Xu and Shen, 2015; Xu and Ma, 2018) and some ginglymodians (Bürgin, 2004; Tintori and Lombardo, 2007; López-Arbarello et al., 2011; Xu and Wu, 2012; Xu et al., 2018). Up to date, 14 holostean species (in 13 genera) have been reported from the Middle Triassic marine rock succession in Yunnan and Guizhou provinces, Southwest China (Xu, 2019). They are important for our understanding on the early evolution of holosteans, but some taxa are poorly known in morphology and controversy in taxonomy, urgently needing detailed description and revision.

Sinoeugnathus kueichowensis Su, 1959 is such a taxonomically controversial holostean that was incompletely described on the basis of a single specimen (IVPP V2443) from the Middle Triassic (Ladinian) marine deposits exposed in the Dingxiao, Xingyi, Guizhou Province. Su (1959) regarded *Sinoeugnathus* as a close relative of *Eoeugnathus* from the Middle Triassic of Italy and Switzerland (Brough, 1939; Herzog, 2003), and classified it within the halecomorph family Eugnathidae Nicholson & Lydekker, 1889, a junior synonym of Caturidae Owen, 1860 (Patterson, 1973). Unfortunately, the holotype of *S. kueichowensis* was later missing for unknown reason, and our understanding of this taxon has long been provided only by the original description and illustration of Su (1959). In the last decade, 15 well-preserved specimens that can be referred to this species were collected by the second author at the same fossil level from Xingyi in Guizhou, and Fuyuan and Shizong in Yunnan. Studies of these specimens showed that some morphological details were previously undescribed or misidentified for *Sinoeugnathus*, and accordingly the attribution of this taxon to the Caturidae is now in doubt. A taxonomic revision of this taxon is present in this paper.

Sinoeugnathus kueichowensis was recovered from dark grey thin- to medium-bedded marlites and argillaceous limestones at the lower part of the Zhuganpo Member of the Falang Formation in western Guizhou and eastern Yunnan. Besides *Sinoeugnathus*, the coexisting vertebrates include several other types of ray-finned fishes (Su, 1959; Jin, 2001; Liu et al., 2002, 2003; Xu et al., 2012, 2015, 2018; Xu and Ma, 2018; Tintori et al., 2014, 2015; Xu, 2020), a coelacanth (Geng et al., 2009), and diverse marine reptiles (Young, 1958; Li, 2006; Li C et al., 2016; Jiang et al., 2020; Shang et al., 2020). The whole fossil assemblage was named the Xingyi Fauna or Biota (Li, 2006; Benton et al., 2013). Biostratigraphical studies of ammonoids (Zou et al., 2015) suggest a late Middle Triassic (Ladinian) age for this fauna, and this age determination is supported by a zircon U-Pb dating (240.8 ± 1.8 Ma) for the fossil beds (Li et al., 2016).

2 Material and methods

All specimens are curated at the fossil collections of the Institute of Vertebrate Paleontology and Paleoanthropology (IVPP), Chinese Academy of Sciences. They were mechanically prepared with sharp steel needles. For better contrast, some specimens were dusted with ammonium chloride (NH_4Cl) before being photographed. The relative position of fins and scale counts were expressed following Westoll (1944). To illuminate the phylogenetic position of *Sinoeugnathus* within the Holostei, we incorporated it into a data matrix slightly expanded from Xu (2019). A new character was proposed (Char. 225 in the electronic supplementary material), and the current data matrix includes 225 morphological characters coded across 62 extant and fossil terminal taxa. All characters were unordered and equally weighted. Besides *Sinoeugnathus*, two taxonomically controversial halecomorph taxa, *Eoeugnathus* and *Allolepidotus* from the Middle Triassic of Europe (Brough, 1939; Lombardo, 2001; Herzog, 2003), were added as well. *Pteronisculus stensiöi* was selected for out-group comparison. Tree searches were accomplished with the heuristic search algorithm in PAUP* (v. 4.0a169) using 1000 random addition sequence replicates, holding five trees at each step, with the tree bisection and reconnection (TBR) strategy enabled and maxtrees set to automatically increase by 100 (Swofford, 2003).

Anatomical abbreviations ang, angular; ao, antorbital; apl, anterior pit line; bf, basal fulcrum; br, branchiostegal ray; cl, cleithrum; den, dentary; dpt, dermopterotic; dsp, dermosphenotic; es, extrascapular; ff, fringing fulcrum; fr, frontal; gu, gular; hm, hyomandibula; hmf, foramen for the hyomandibular trunk of facial nerve (VII); io, infraorbital; iop, interopercle; le, lateral ethmoid; mpl, middle pit line; mx, maxilla; na, nasal; op, opercle; pa, parietal; pas, parasphenoid; pcl, postcleithrum; pf, principle fin ray; pmx, premaxilla; pop, preopercle; ppl, posterior pit line; prr, procurent ray; pt, posttemporal; qj, quadratojugal; qu, quadrate; r, rostral; scl, supracleithrum; scr, sclerotic bones; smx, supramaxilla; so, suborbital; sop, subopercle; sp, sphenotic; su, supraorbital; sy, symplectic; vo, vomer.

3 Systematic paleontology

Actinopterygii Cope, 1887

Neopterygii Regan, 1923

Holostei Müller, 1845

Halecomorphi Cope, 1872

Ionoscopiformes Grande & Bemis, 1998

Subortichthyidae fam. nov.

Diagnosis Small-sized ionoscopiforms distinguished from other members of this order by the following derived features (those unique among ionoscopiforms identified with an asterisk): frontal about four times as long as parietal; two or three supraorbitals; one to three suborbitals; quadrate almost fully covered by posterior portion of maxilla (*); elongate maxilla

extending posterior to coronoid process of lower jaw (*); supramaxillary process of maxilla relatively small; 12 pairs of branchiostegal rays; 30–35 lateral line scales (*); complete row of elongate scales between last lateral line scale and uppermost caudal fin ray (*).

Included genera *Subortichthys*, *Sinoeugnathus*, *Eoeugnathus* and *Allolepidotus*.

Type genus *Subortichthys*.

Geographical distribution and age Luoping, Fuyuan and Shizong, Yunnan, and Xingyi, Guizhou, China; Lombardy, Italy; Grisons and Canton Ticino, Switzerland. Pelsonian (Anisian) to Ladinian, Middle Triassic.

Sinoeugnathus kueichowensis Su, 1959

Neotype IVPP V24315, a nearly complete, lateral compressed specimen with a standard length of 64.2 mm from Xiemi, Wusha, Xingyi.

Referred specimens IVPP V19007, 30793 and 30794 from Xiemi Village, Wusha Town, Xingyi City, Guizhou Province; V24314, 24316, 24317, 24320, 30784–30787 from Jiyangshan Village, Fuyuan County, Yunnan Province; V30788–30790 from Shizong County, Qujing City, Yunnan Province.

Locality and horizon Xiemi and Wusha, Guizhou; Jiyangshan, Fuyuan and Shizong, Yunnan. Zhuganpo member of the Falang Formation, Ladinian, Middle Triassic.

Emended diagnosis A subortichthyid distinguished from other members of this family by the following features (those unique among subortichthyids identified with an asterisk): parietal rectangular, with width/length ratio of ~ 0.7 ; dermopterotic nearly twice as long as parietal; antorbital hatchet-shaped, with single large sensory pore at its posterior portion (*); three pairs of extrascapulars; third infraorbital relatively large but free from contact with preopercle; two large suborbitals extending posteriorly and partly covering preopercle; opercle fan-shaped, with length nearly equal to depth; 15 rays in each pectoral fin; seven principal dorsal rays; nine principal anal rays; 15 principal caudal fin rays; and scale formula of D16/P6–7, A14, C28–29/T33–34.

4 Comparative description

General morphology and size *Sinoeugnathus kueichowensis* is a small-sized ionoscopiform with a blunt snout, a fusiform body and a moderately forked caudal fin. The dorsal fin originates slightly behind the pelvic fins, and the anal fin inserts at the posterior 1/4 of the body. Among ten nearly completely preserved specimens, the standard lengths range from 52.2 to 69.6 mm (Table 1). The maximum body depth lies midway between the posterior margin of the opercle and the origin of the dorsal fin. In the neotype (Fig. 1A), the head length and body depth account for 33% and 42% of the standard length, respectively. The outer surfaces of cranial bones are ornamented with ganoine tubercles and ridges (Figs. 2–4). The general body form (Fig. 1C) can be reconstructed based on V24315 (Fig. 1A), and V30784 (Fig. 1B).

Snout The canal-bearing bones in the snout region consist of a median rostral and paired nasals and antorbitals (Figs. 2–4). The rostral is small and curved, with a concave

anterior margin and a convex posterior margin. It contacts the nasal dorsally, the antorbital laterally, and the premaxilla ventrally. The anterior commissure of the lateral line system is enclosed in this bone (Figs. 2–4). The nasals are elongate with a tapering posterodorsal tip inserting into the anterior portion of the frontal. The lateral margin of the nasal is concaved for the posterior nostril. Analogous to the living bowfin, the anterior nostril is probably located between the nasal, rostral and antorbital. The antorbitals are hatchet-shaped, having an elongate anteroventral extension and an expanded posterior portion. Each antorbital has an anterior contact with the rostral and premaxilla, and posteriorly forms a part of the anterior orbital margin. A large sensory pore is discernable at the posterior portion of this bone (Figs. 2–4).

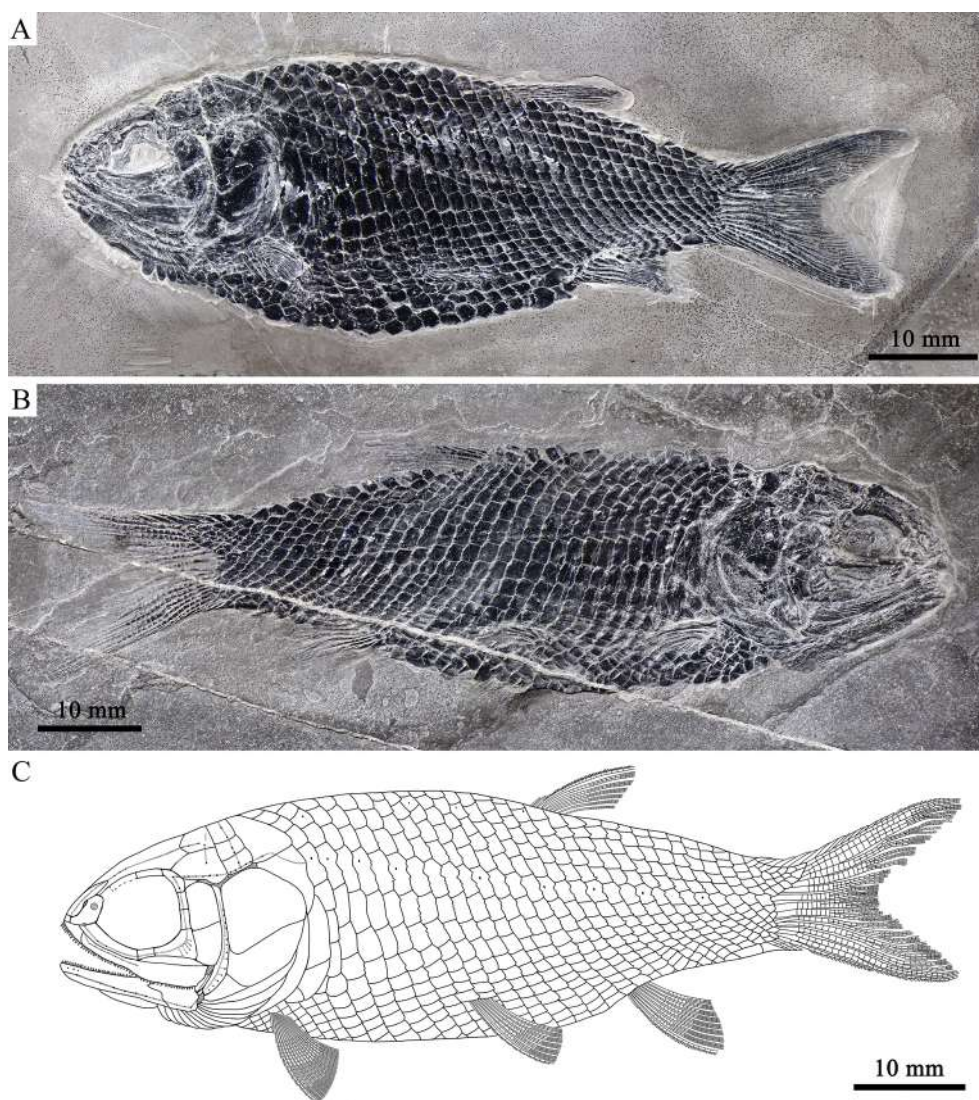


Fig. 1 Two complete specimens and reconstruction of *Sinoeugnathus kueichowensis*
A. IVPP V24315 (neotype); B. V30784; C. reconstruction

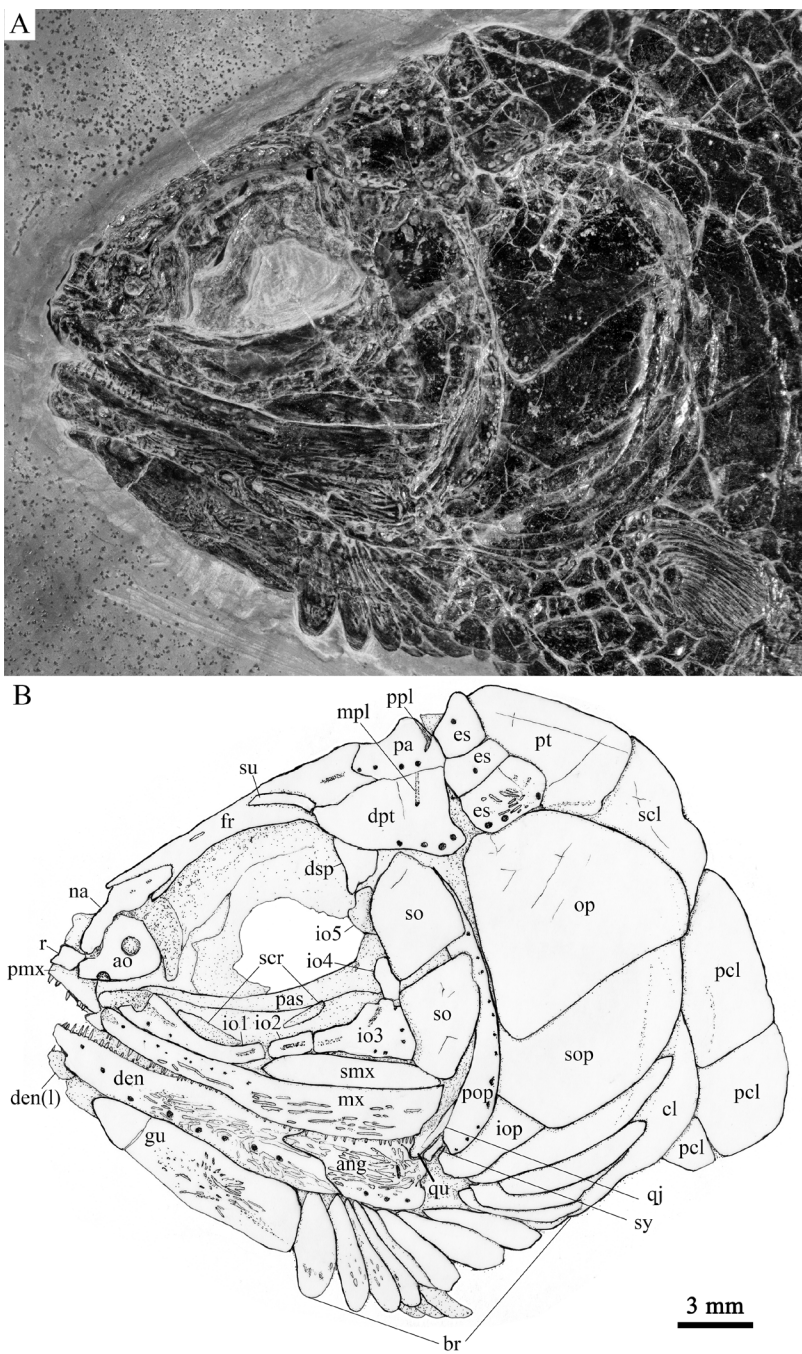


Fig. 2 Photograph (A) and line-drawing (B) of skull and pectoral girdle of *Sinoeugnathus kueichowensis*, IVPP V24315 (neotype)

Skull roof The skull table consists of a pair of frontals, parietals, dermopterotic and three pairs of extrascapulars. The frontal, the largest component of the skull roof, is slightly more than four times as long as the parietal. It is roughly trapezoid, slightly constricted above

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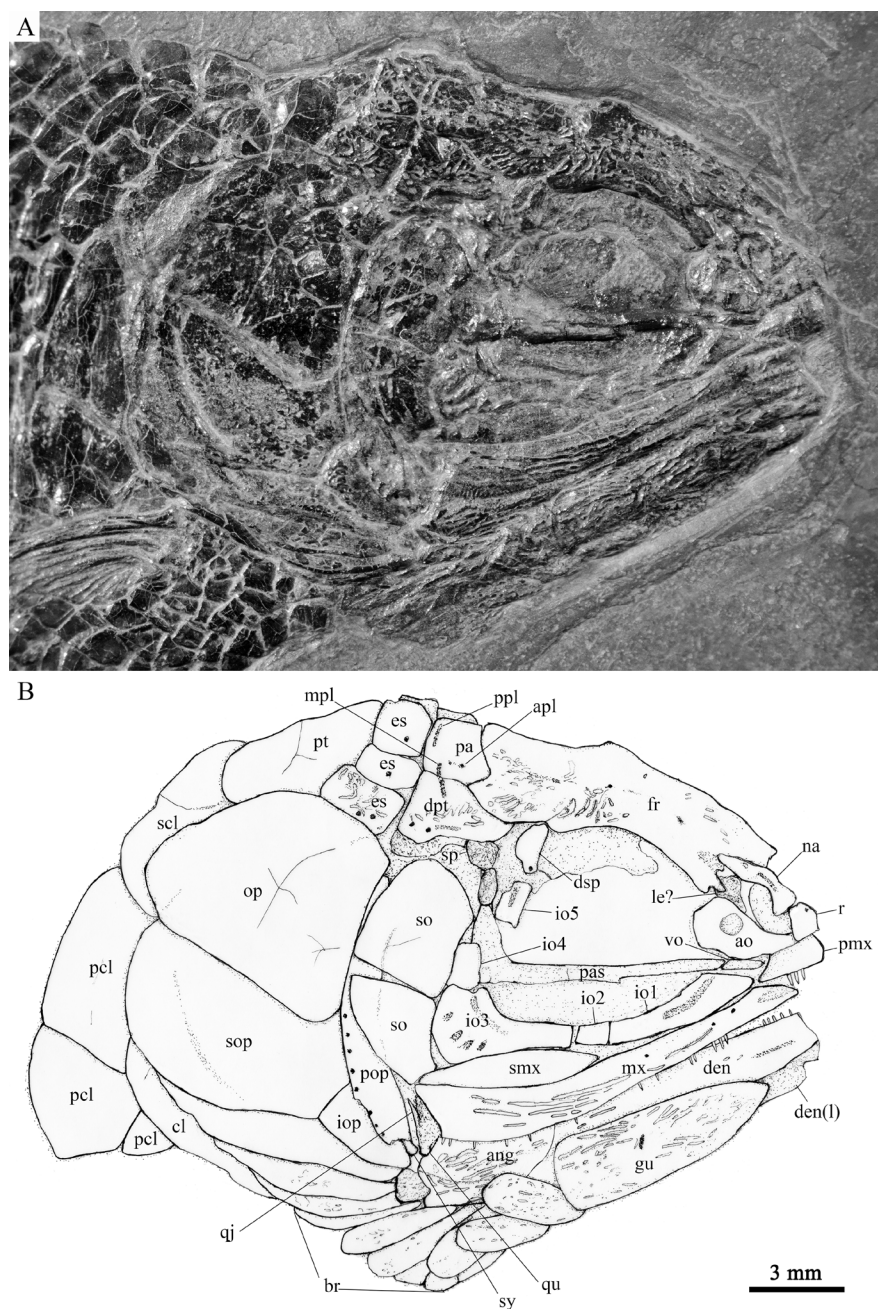


Fig. 3 Photograph (A) and line-drawing (B) of skull and pectoral girdle of *Sinoeugnathus kueichowensis*, IVPP V30784

the orbital region. The median suture between frontals is nearly straight. The anterior margin is concave, receiving the posterior portion of the nasal (Figs. 2–4). The supraorbital sensory canal enters this bone from the nasal, runs longitudinally through the bone and enters the parietal posteriorly. The parietal is small and nearly rectangular, with a width/length ratio of

about 0.7. It has a straight lateral margin bordering the dermopterotic and a convex anterior margin bordering the frontal. Three pit-lines originate in this bone (Fig. 3). The anterior pit-line is situated at the anterolateral portion of the parietal. The middle pit-line originates at the posterolateral portion of the bone and extends laterally into the dermopterotic. The posterior pit-line is located at the posterior portion of the parietal, extending medially for a short length in this bone.

The dermopterotic is trapezoidal, about twice as long as the parietal. It has a tapered anterior process that fits a posterior notch of the frontal (Figs. 2–4). The temporal sensory canal extends longitudinally through this bone and enters the extrascapular posteriorly, indicated by a series of pores parallel to the lateral margin of this bone (Figs. 2, 3). Three extrascapulars are nearly trapezoidal or rectangular and their sizes slightly vary in different specimens. The supratemporal commissure runs transversely through the extrascapulars, indicated by several pores at the middle portions of these bones.

Circumorbital bones There are at least one supraorbital and five infraorbitals. A slightly detached, posterior supraorbital is discernable in V24315 (Fig. 2). It is elongate and trapezoidal, being about one-third of the orbital length. The first (anterior-most) infraorbital is long and cleaver-shaped, slightly expanded anteriorly. The second infraorbital is short and rectangular. The third infraorbital is large and pentagonal, tapering anteriorly. The fourth and fifth infraorbitals are small and slightly deeper than long, contacting the suborbitals posteriorly, the dermosphenotic dorsally and the third infraorbital ventrally. The infraorbital sensory canal extends through the infraorbitals, indicated by some pits and canals near the orbital margins of these bones (Fig. 2).

The dermosphenotic is trapezoidal, similar to the last infraorbital in size (Figs. 2–4). It narrows ventrally and receives the infraorbital sensory canal from the latter bone. The sphenotic, unfused with the dermosphenotic, has an exposed dermal component on the skull roof as commonly in other holosteans.

Two suborbitals are present. Both are broad and trapezoidal; the dorsal is larger, deeper than wide, and the ventral is relatively small, tapered ventrally (Figs. 2–4).

Two sclerotic bones are partly discernable near the orbit rim in V24315 (Fig. 2). They are thin and slightly curved.

Parasphenoid, palatoquadrate and suspensorium The parasphenoid and palatoquadrate can only be discerned through the orbit laterally; the former is elongate, contacting the latter laterally (Fig. 2). The quadrate is only partly exposed, with its anterodorsal portion laterally covered by the maxilla. It bears a strong condyle that articulates the lower jaw. The quadratojugal is narrow and splint-like (Figs. 2–4). It articulates with the posterolateral surface of the ventral portion of the quadrate, and resets on the anterior edge of the preopercle.

The hyomandibula is laterally covered by two suborbitals and the preopercle and is not exposed in the specimens. X-ray scanning of V30785 reveals a hatchet-shaped and vertically positioned hyomandibula, with a foramen for hyomandibular branch of the facial nerve in the

middle. It has a single dorsal facet articulating the posteroventral side of the brain case (Fig. 5D).

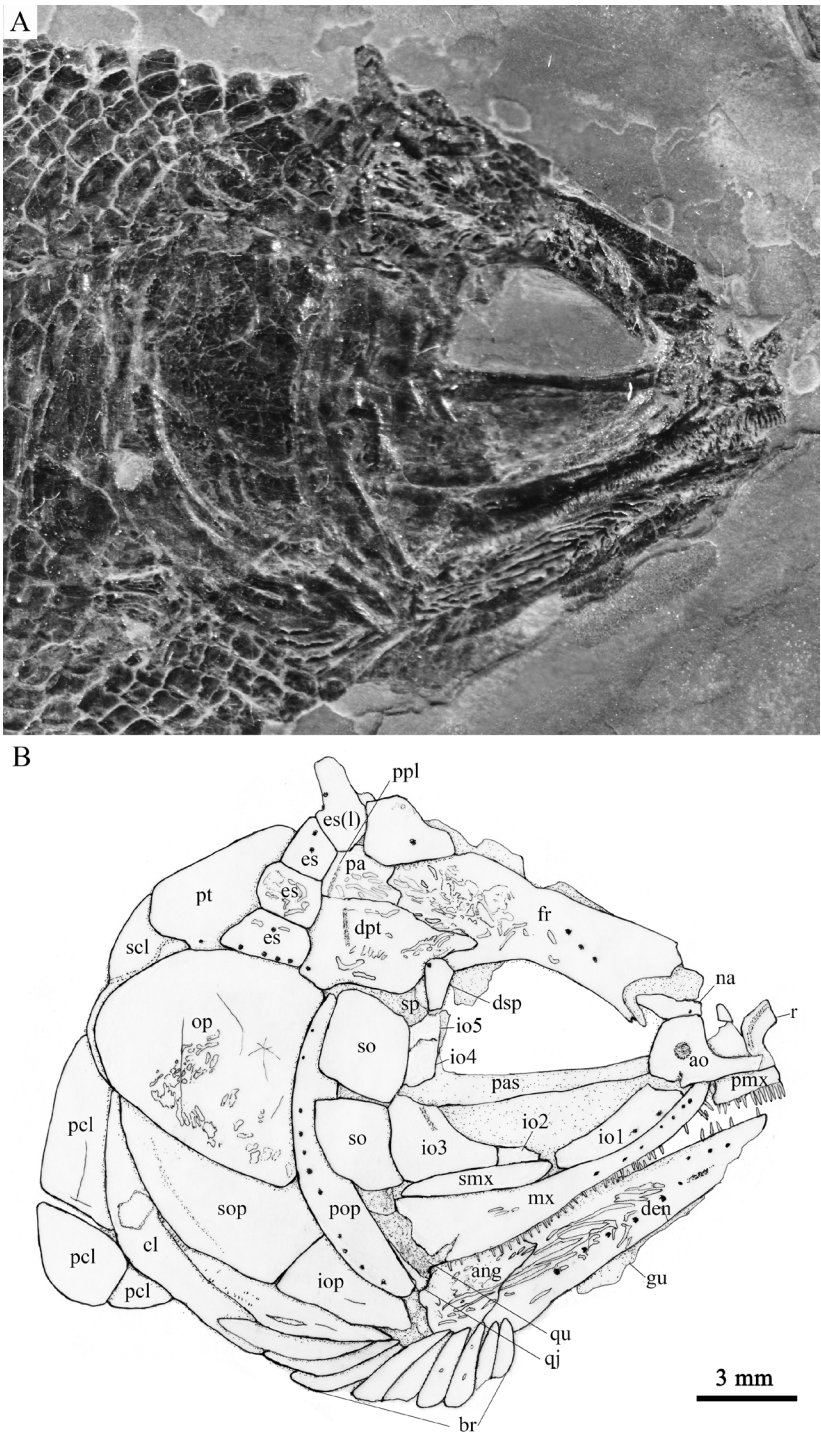


Fig. 4 Photograph (A) and line-drawing (B) of skull and pectoral girdle of *Sinoeugnathus kueichowensis*, IVPP V30785

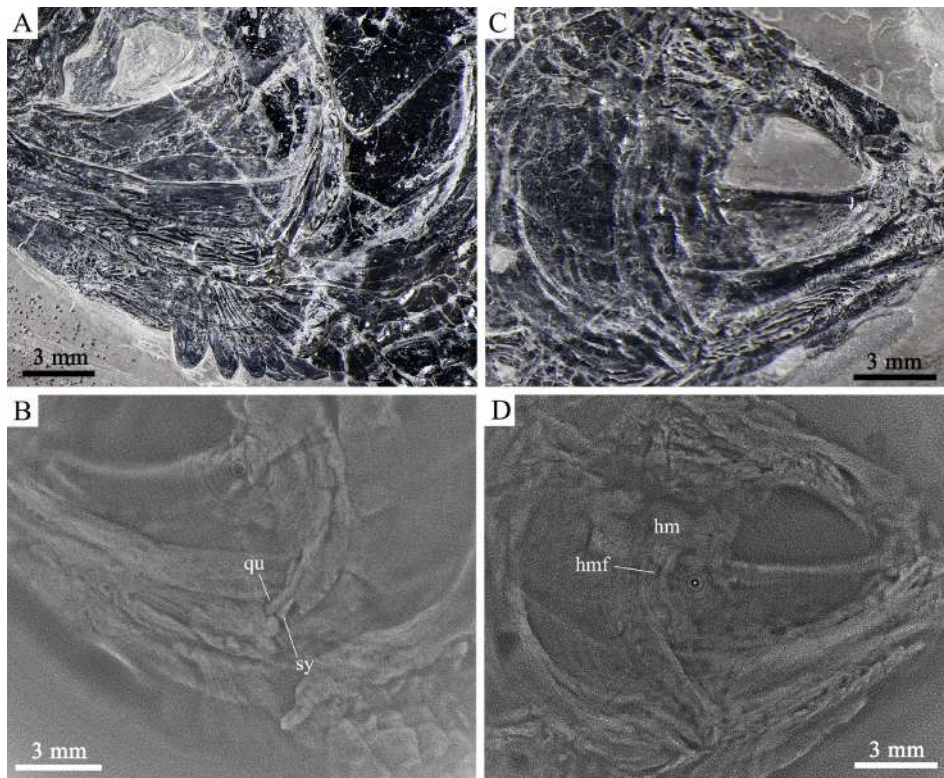


Fig. 5 Cranial bones of *Sinoeugnathus kueichowensis*

A. IVPP V24315; B. X-ray computed tomography of A, showing both the quadrate symplectic condyles for the jaw hinge; C. V30785; D. X-ray computed tomography of C, showing the hyomandibula and the foramen for the hyomandibular trunk of facial nerve (VII)

Most of the symplectic is laterally covered by the preopercle except for the ventral condyle of this bone. This condyle, together with the quadrate condyle, forms a double jaw joint with the lower jaw (Figs. 2, 3, 5D).

Jaws The premaxilla is relatively large, having a horizontally expanded oral region and a posterodorsally directed nasal process (Figs. 2–4). It is uncertain whether the foramen for the olfactory nerve is present in the nasal process of this bone or not because of the overlapping of nasals. The maxilla is elongate, ending at the level of the jaw articulation. The middle portion of this bone slightly protrudes dorsally and forms a rudimentary supramaxillary process. Posterior to this process, the dorsal margin of the maxilla is distinctly concave for accommodating a supramaxilla (Figs. 2–4). The posterior margin of the maxilla is slightly concave, resembling the condition in *Subortichthys*. As in other ionoscopiforms, the maxilla encloses a branch of the infraorbital sensory, indicated by a longitudinal row of pores at the anterior half of the maxilla. The supramaxilla is about half of the maxillary length. It is elongate, tapering at both ends. The oral margin of the maxilla is fully equipped with small conical teeth, over forty in number (Fig. 2). Additionally, a row of 12 conical teeth is

discernable along the oral margin of the premaxilla and they are slightly larger than those on the anterior portion of the maxilla (Fig. 3).

The lower jaw is elongate and not strong, having a height/length ratio of 0.18. The dentary is wedge-shaped, being the largest element of the lower jaw. It contacts the relatively small, trapezoidal angular posteriorly in a sinuous suture. The mandibular sensory canal extends longitudinally through the dentary and angular, indicated by eight or nine pores in the former and three pores in the latter (Figs. 2, 4). Three coronoid bones are exposed medially to the dentary (V30791). They are covered with small conical teeth.

Opercular series The preopercle is narrow and crescent-shaped with part of its dorsal portion anteriorly covered by two suborbitals. The dorsal tip of the preopercle contacts the posterolateral process of the dermopteroic and its anteroventral tip nearly reaches the ventral end of the quadratojugal. The preopercular sensory canal runs dorsoventrally through the preopercle. Additionally, a series of pores near the posterior margin of this bone likely represents the openings for the branches of the preopercular sensory canal (Figs. 2–4). The opercle is large and fan-shaped, with a depth nearly equal to its length. The posterior and dorsal margins are rounded and the anterior and ventral margins are straight (Figs. 2–4). The subopercle is relatively small and nearly sickle-shaped, bearing a short triangular anterodorsal process. The interopercle is small and triangular. It tapers anteroventrally with an anterodorsal portion laterally covered by the preopercle (Figs. 2–4).

Gular and branchiostegal rays The median gular is elongate and sub-circular, with a pointed anterior tip and a relatively broad posterior end (Figs. 2, 3). Its length is slightly over half length of the lower jaw. Twelve pairs of branchiostegal rays are present in V24315 and V30784 (Figs. 2, 3), representing the maximum number in this taxon. They are elongate and plate-like, increasing in length posteriorly. The median gular and branchiostegal rays are ornamented with some tubercles and ridges.

Girdles and fins A posttemporal, a supracleithrum, a cleithrum and three postcleithra are present at each side of the pectoral girdle. The posttemporal is sub-triangular, with a rounded posterior margin. It tapers medially with a narrow anterior portion overlapped by the extrascapulars. The lateral line extends longitudinally through the lateral portion of posttemporal, and enters the supracleithrum (Fig. 2). The supracleithrum is deep and anteriorly inclined, with its anterior portion slightly covered by the posttemporal and opercle. The complete shape of the cleithrum remains unknown because of overlapping of the opercular series *in situ* (Figs. 2–4). The exposed portion of the cleithrum is largely crescent-shaped with its dorsal arm nearly equal to the horizontal arm (Figs. 2–4). There are three postcleithra associated with the cleithrum. The dorsal is the largest, as deep as the supracleithrum; the middle is trapezoidal, near half of the size of the dorsal; and the ventral is smallest and triangular (Figs. 2–4).

The relatively large pectoral fins insert low on the body and each bear 15 distally segmented rays. The first, leading ray is unbranched, preceded by a basal fulcrum and a series

of fringing fulcra. The remaining rays are distally segmented (Fig. 6A).

The pelvic girdles are not exposed. The small pelvic fins insert at the 6th or 7th vertical scale row, and each consists of seven distally segmented rays. Several fringing fulcra are associated with the leading ray of the pelvic fin (Fig. 6B).

Median fins The dorsal fin originates at the 16th vertical scale row, composed of seven distally segmented rays. The first ray is unbranched, preceded by two to three basal fulcra and several fringing fulcra; the remaining rays are branched distally (Fig. 6C).

The anal fin originates below the 14th vertical scale row, having nine distally segmented principal rays. The first principal ray is unbranched, preceded by a short rudimentary ray, a basal fulcrum and a series of fringing fulcra, and the remaining rays are branched distally. The rudimentary ray is composed of two segments, less than one-fourth of the length of the first principal ray (Fig. 6E).

The caudal fin is hemi-heterocercal, with a moderately forked profile. It has 15 principal rays, seven in dorsal lobe. The dorsal and ventral marginal principal rays are segmented and unbranched, and the middle principal rays are segmented and branched up to three times. There are seven basal fulcra in the dorsal lobe and one or two basal fulcra and three rudimentary rays in the ventral lobe. Small leaf-like fringing fulcra are present in both lobes (Fig. 6D).

Squamation The body is fully covered with rhomboid scales (Figs. 1, 6, 7). The scales are arranged in 33–34 vertical rows along the main lateral line. In addition, five rows of scales extend into the epaxial lobe of the caudal fin. Besides the ridge scales, there are 15–17 scales in the tenth vertical row on each side of body, ten below the lateral line. The scales in the anterior flank region are about three times deeper than wide, and they gradually become shorter and smaller dorsally, ventrally and posteriorly. Moreover, an enlarged lateral scute is discernable anterior to the anal fin. The lateral line scales bear a distinct notch in the middle of its posterior margin, which is likely for the opening of the lateral line. An additional lateral line is present, indicated by a series of small pores in the predorsal region. Most of the scales, except those covering the epaxial lobe of the caudal fin, have a serrated posterior margin with 1–4 acute projections. The external surfaces of scales are largely smooth except for small pores on some scales.

Table 1 Measurement data of ten specimens of *Sinoeugnathus kueichowensis* (mm)

Specimen	SL	HL	BD	TL	PVL	PDL	PAL
V19007	66.20	20.88	23.16	83.28	–	40.16	52.24
V24315	64.22	20.94	26.92	82.56	35.18	41.70	49.66
V24316	52.20	18.98	22.48	63.68	29.00	29.28	40.32
V24317	58.66	20.40	22.24	73.06	33.92	37.90	45.10
V24320	65.40	22.74	25.00	83.06	–	40.14	49.24
V30784	69.58	22.12	23.42	87.54	37.44	39.52	52.24
V30786	59.68	19.06	22.86	70.78	34.16	37.46	46.38
V30787	61.32	21.18	22.36	69.12	31.52	35.98	44.14
V30789	60.50	22.04	21.80	75.66	31.32	–	44.46
V30790	56.56	21.44	21.72	74.88	30.50	37.52	42.92

Abbreviations: BD, body depth; HL, head length; PAL, preanal length; PDL, predorsal length; PVL, prepelvic length; SL, standard length; TL, total length.

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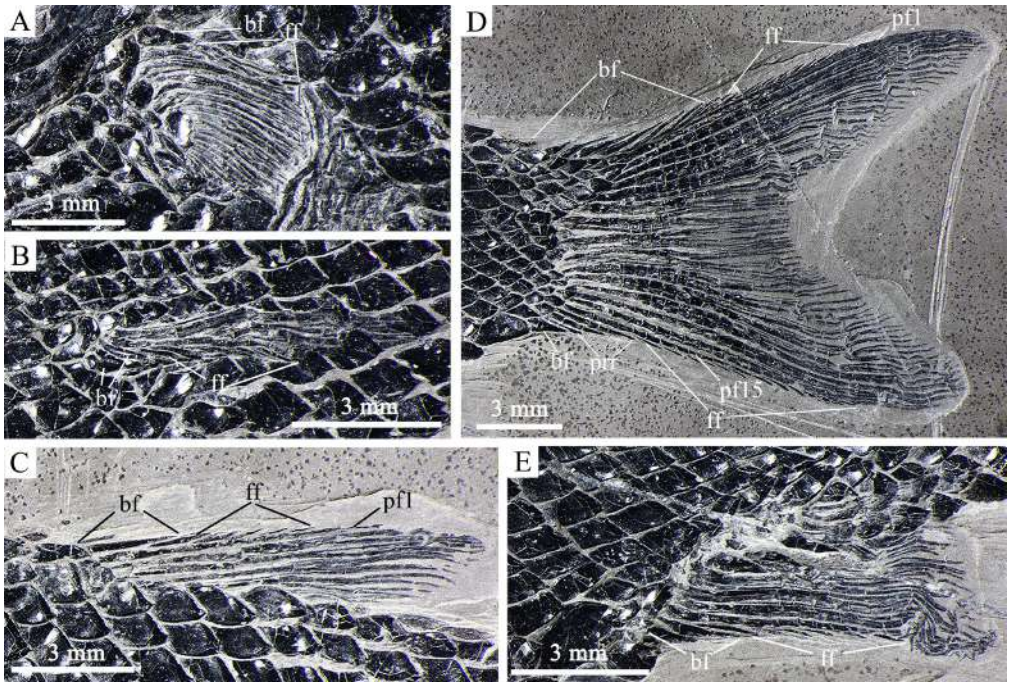


Fig. 6 Fins of *Sinoeugnathus kueichowensis*, IVPP V24315
A. left pectoral fin; B. left pelvic fin; C. dorsal fin; D. caudal fin; E. anal fin

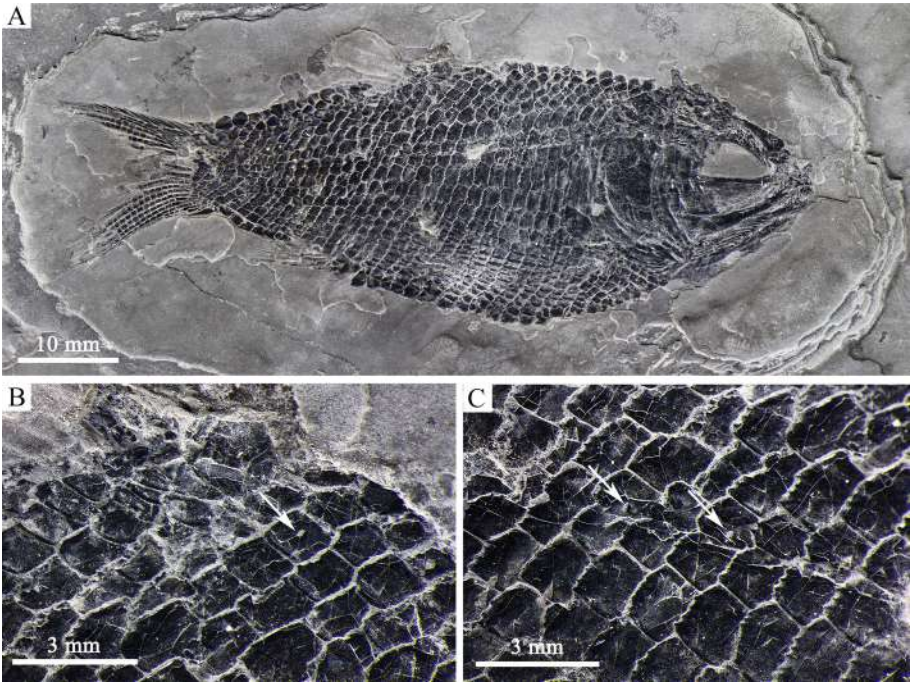


Fig. 7 Complete specimen and lateral line scales of *Sinoeugnathus kueichowensis*, IVPP V30785
A. complete specimen; B. scales in predorsal region with arrow showing pore of additional lateral line;
C. main lateral line scales in posterior flank region with arrows showing pores of lateral line

5 Discussion

5.1 New specimens and neotype

The new specimens are referred to *Sinoeugnathus kueichowensis* based on their similarity to the missing holotype and previously only known specimen illustrated by Su (1959) in morphology and size. Most strikingly, the maxillae in the new specimens are quite long, accounting slightly over half of the head length. Such a long maxilla, also present in the holotype of *S. kueichowensis*, has not known in other holosteans from the same fossil beds (Liu et al., 2002, 2003; Xu and Ma, 2018; Xu et al., 2018). Additionally, the opercle in the new specimens is large and fan-shaped with rounded posterior and dorsal margins, consistent with the condition in the illustrated holotype (Su, 1959). Moreover, the new specimens and holotype show the same scale formula (D17/P6, A14, C27/T33). The proportions of the head length and body depth to the standard length in the new specimens (Table 1) are very similar to those of the holotype (SL = 53 mm; HL = 19 mm; BD = 22 mm). Among the new specimens, IVPP V24315 is best-preserved and relatively large, and is therefore designated as the neotype of *S. kueichowensis*.

5.2 Phylogenetic analysis

Phylogenetic analysis recovered 48 most parsimonious trees (Tree length = 661 steps; consistency index = 0.4115; retention index = 0.7665). In the strict consensus tree (Fig. 8), Parasemionotiformes (sampled by *Watsonulus*) is nested at the base of the Halecomorphi, followed successively by Ionoscopiformes and Amiiformes, consistent with previous studies (Grande and Bemis, 1998; Xu and Ma, 2018; Xu, 2019). *Sinoeugnathus* possesses two synapomorphies of the Halecomorphi: a symplectic articulating with the lower jaw, and a notched posterior margin of the maxilla. Within this clade, it is resolved as an ionoscopiform, possessing a synapomorphy of this order, a sensory canal in the maxilla (Grande and Bemis, 1998; Xu, 2019). *Sinoeugnathus* is sister to the Anisian ionoscopiform *Subortichthys* from Luoping, Yunnan, and both, together with two European genera *Allolepidotus* and *Eoeugnathus*, form a monophyletic group (Subortichthyidae fam. nov.) at the base of Ionoscopiformes.

5.3 Taxonomic revision and comparison

Su (1959) noticed the resemblances of *Sinoeugnathus* to *Eoeugnathus* in general body form, maxilla and opercle, and referred the genus to the Eugnathidae (= Caturidae; Patterson, 1973). Besides these two genera, *Allolepidotus* and several other Triassic genera (e.g., *Heterolepidotus* and *Furo*) were previously also included in this family (Gardiner, 1960; Lombardo, 2001), but this taxonomic assignment was questioned by Patterson (1973), who placed three ‘caturids’ (*Allolepidotus*, *Eoeugnathus* and *Sinoeugnathus*) into the Parasemionotidae. However, parasemionotids are now confined to basal halecomorphs in the Early Triassic (Grande and Bemis, 1998). Later, Herzog (2003) made a revision of

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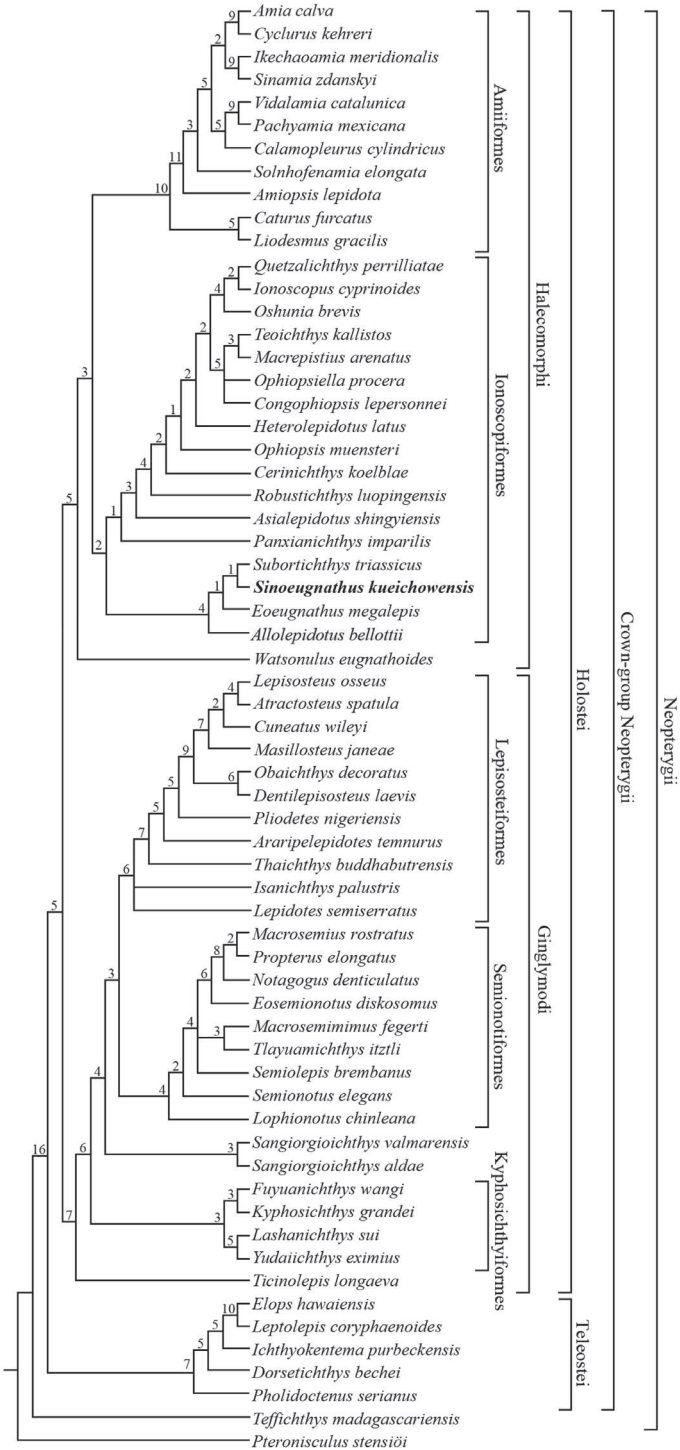


Fig. 8 Strict consensus of 48 most parsimonious trees illustrating the phylogenetic relationships of *Sinoeugnathus* within the Neopterygii

Tree length = 661 steps; consistency index (CI) = 0.4115; retention index (RI) = 0.7665

Digits above nodes indicate Bremer decay indices. For character descriptions and codings for the sampled taxa, see the online supplementary material

Eoeugnathus and placed it into the Section B (the clade Ionoscopiformes plus Amiiformes) of the Halecomorphi without a reference to a special family or order. Ebert (2018) incorporated *Eoeugnathus* in a phylogenetic analysis, and recovered it as a sister taxon of *Allolepidotus* within ‘Panxianichthyiformes’ (Sun et al., 2016). However, the grade ‘Panxianichthyiformes’ is likely paraphyletic, comprising basal members of Ionoscopiformes (Xu and Ma, 2018; Xu, 2019).

Our study confirms that *Sinoeugnathus* is not a caturid as previously suggested (Su, 1959) because it lacks diagnostic features of this family, e.g., sharply carinate acrodin tooth caps on larger jaw teeth, an extremely slender rod-like maxilla, and 22 or more pairs of branchiostegal rays (López-Arbarello and Ebert, 2023). Instead, this genus is revised here as a subortichthyid, sharing four derived features with other members of this family (*Allolepidotus*, *Eoeugnathus* and *Subortichthys*): a quadrate almost fully covered by the maxilla; a maxilla extending posterior to the coronoid process of the lower jaw; presence of 35 or less lateral line scales; and a complete row of elongate scales between the last lateral line scale and the uppermost caudal fin ray. Although these features have independently evolved in some other neopterygians, they are absent in other ionoscopiforms.

Sinoeugnathus is easily distinguished from other members of this family by the following features: 1) *Sinoeugnathus*, similar to *Subortichthys* (Ma and Xu, 2017), has three pairs of extrascapulars (Figs. 2–4), but *Eoeugnathus* (Herzog, 2003) and *Allolepidotus* (Lombardo, 2001) have only single pair of extrascapulars; outside of Subortichthyidae, three pairs of extrascapulars have independently evolved in the ionoscopiform *Aisalepidotus* (Xu and Ma, 2018); 2) *Sinoeugnathus* has two suborbitals (Figs. 2–4), contrasting the counts in other subortichthyids (single in *Allolepidotus*; and three in *Subortichthys* and *Eoeugnathus*); 3) the antorbital in *Sinoeugnathus* is relatively short and deep (Figs. 2–4), but the bone in other subortichthyids is relatively long and slender (Ma and Xu, 2017; Lombardo, 2001; Herzog, 2003); 4) *Sinoeugnathus* has a fan-shaped opercle with a depth nearly equal to its length (Figs. 2–4); by contrast, the opercle in other subortichthyids is commonly trapezoidal, deeper than long; 5) *Sinoeugnathus* has only 15 principal caudal fin rays (Fig. 6D), fewer than those in other subortichthyids (18 in *Subortichthys* and *Eoeugnathus*, and ~20 in *Allolepidotus*); and 6) the lateral line scales in the anterior flank region of *Sinoeugnathus* are notably deep (nearly 3 times deeper than wide), but those in other subortichthyids are relatively short (1.2–2 times deeper than wide).

5.4 Ecological implications

Subortichthyids generally have a maximum standard length no more than 100 mm. Among this family, *Sinoeugnathus* is the smallest one with a maximum standard length of 69.6 mm, and other subortichthyids are slightly larger (SL = 72 mm in *Subortichthys*, 85 mm in *Allolepidotus* and 98 mm in *Eoeugnathus*). By contrast, other Triassic ionoscopiforms are significantly larger (SL = 159 mm in *Panxianichthys*, 273 mm in *Aisalepidotus*, and 360

mm in *Robustichthys*; Table 2). Additionally, a difference between subortichthyids and other Triassic ionoscopiforms is about the dentation. The former has smaller and more numerous conical teeth in the relatively slender and longer jaws than the latter. Moreover, subortichthyids have a shorter coronoid process of the lower jaw with higher ratios of mandibular length to skull length and of maxillary length to mandibular length than those of non-subortichthyid ionoscopiforms (Table 2). Notably, the teeth on the palatine are very weak in subortichthyids, but they are stronger and blunt in other Triassic ionoscopiforms (e.g., *Asialepidotus*; Xu and Ma, 2018). Analogous to modern ray-finned fishes, the morphological divergence between small-sized subortichthyids and larger non-subortichthyid ionoscopiforms was probably driven by ecological specializations related to feeding and foraging, e.g., the type of preys they consumed, the reaction distance and swimming performance (Folkvord and Hunter, 1986; Scharf et al., 2000; Magnhagen and Heibo, 2001; Karpouzi and Stergiou, 2003). The absence of blunt palatine teeth indicates that subortichthyids probably fed on relatively soft preys mainly by biting (rather than by crushing). As one of the small-sized holosteans in the Middle Triassic Xingyi Biota, *Sinoeugnathus* likely fed on some small shrimps or other relatively soft invertebrates. It was probably in turn fed by larger carnivorous fishes or marine reptiles in the same marine ecosystem (Benton et al., 2013).

Subortichthyids represent an early radiation of ionoscopiforms from the Middle Triassic of both Europe and South China, which at then were located at the eastern and western realms of the Paleo-Tethys Ocean, respectively (Metcalfe, 2011). The recovery of European subortichthyids (*Allolepidotus* and *Eoeugnathus*) as successive sister taxa to Chinese subortichthyids suggests that this family might originate in the western Tethys realm, and the Paleo-Tethys Ocean would provide a west–east corridor for the long dispersal of subortichthyids into the eastern Tethys realm. This paleobiogeographic scenario contradicts the current stratigraphic records of subortichthyids; the European subortichthyids are Ladinian, nearly coeval with *Sinoeugnathus* but slightly younger than the Anisian *Subortichthys* (the oldest record of this family from China). However, this contradiction was possibly caused by incomplete fossil record of subortichthyids from Europe. If our current scenario of subortichthyid phylogeny is accepted, Anisian or even older record of subortichthyids are expected to be found in the western Tethys realm.

Table 2 Measurements for eight specimens of Triassic ionoscopiforms (mm)

Species	MXL	MDL	HL	MXL/MDL	MDL/HL	SL	Specimen
<i>Sinoeugnathus kueichowensis</i>	13	15	22	0.87	0.68	69.6	IVPP V30784
<i>Subortichthys triassicus</i>	17	19	28	0.89	0.68	72	IVPP V20680
<i>Allolepidotus bellotti</i>	15.5	16.5	26	0.94	0.63	85	MCSNIO P647
<i>Eoeugnathus megalepis</i>	21	25	37	0.84	0.68	98	PIMUZ A/I 1964
<i>Panxianichthys imparilis</i>	20	27	50	0.74	0.54	159	IVPP V19971
<i>Asialepidotus shingyiensis</i>	36	48	91	0.75	0.53	273	IVPP V23010
<i>Robustichthys luopingensis</i>	14	24	45	0.62	0.54	140	IVPP V18568
<i>Robustichthys luopingensis</i>	–	53	100	–	0.53	360	IVPP V20414

Abbreviations: HL, head length; MDL, mandibular length; MXL, maxillary length; SL, standard length.

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6 Conclusion

A detailed examination of 15 new specimens from the Middle Triassic (Ladinian) of Yunnan and Guizhou permitted a thorough revision of the anatomy of *Sinoeugnathus kueichowensis*. This revision provides new insights into the anatomical diversity, phylogeny and paleobiogeography of early ionoscopiforms. A phylogenetic analysis incorporating new anatomical data recovered *Sinoeugnathus* as the sister taxon to *Subortichthys* from the Middle Triassic (Anisian) of Yunnan, and both are grouped with two Ladinian genera *Allolepidotus* and *Eoeugnathus* from the Monte San Giorgio area into a monophyletic group (Subortichthyidae) at the base of the Ionoscopiformes. The previous placement of *Sinoeugnathus* into the Eugnathidae (= Caturidae) is firmly rejected. The proposed scenario of subortichthyid phylogeny combined with the paleogeographic data suggest that the family may originate in the western Tethys realm; early Middle Triassic (Anisian) subortichthyids are expected to be recovered in that realm. The morphological divergence between small-sized subortichthyids and larger non-subortichthyid ionoscopiforms is well recognized. With a maximum standard length of 69.6 mm, *S. kueichowensis* represents the smallest subortichthyid which likely fed on some small shrimps or other relatively soft invertebrates, and it was in turn fed by larger carnivorous fishes or marine reptiles in the same marine ecosystem.

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云贵地区中三叠世近鲱形类贵州中华真颌鱼(*Sinoeugnathus kueichowensis*)的分类学修订

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摘要: 小型近鲱形类(Halecomorphi)贵州中华真颌鱼(*Sinoeugnathus kueichowensis*)发现于中国云贵地区中三叠世拉丁期海相地层中, 曾被归入弓鳍鱼目(Amiiformes)真颌鱼科(Eugnathidae)。基于15块新标本的详细观察, 对该属种进行了重新描述和分类学修订。由

于正模标本已遗失, 指定标本IVPP V24315为新模(标准体长64 mm)。对新标本的研究揭示了该属种在早前研究中一些未被描述或识别错误的形态特征, 包括斧状的眶前骨, 两个宽大的次眶骨, 上颌骨具有感觉管和三对额外肩胛骨。首次将中华真颌鱼纳入近鲱形类分支系统学研究中; 结果显示, 中华真颌鱼与云南罗平安尼期的复兴鱼(*Subortichthys*)构成姐妹群, 两者与欧洲圣乔治山地区拉丁期的*Allolepidotus*和始真颌鱼(*Eoeugnathus*)一起组成预言鱼目(Ionoscopiformes)底部的一个单系群(命名为复兴鱼科)。中华真颌鱼的分类学修订为了解预言鱼目的系统发育关系和古地理演化提供了新的信息。

关键词: 云贵地区, 中三叠世, 近鲱形类, 预言鱼目, 骨骼学, 系统发育

References

- Benton M J, Zhang Q Y, Hu S X et al., 2013. Exceptional vertebrate biotas from the Triassic of China, and the expansion of marine ecosystems after the Permo–Triassic mass extinction. *Earth Sci Rev*, 125: 199–243
- Brough J, 1939. The Triassic fishes of Besano, Lombardy. London: British Museum (Natural History). 1–117
- Bürgin T, 2004. *Eosemionotus ceresiensis* sp. nov., a new semionotiform fish (Actinopterygii, Halecostomi) from the Middle Triassic of Monte San Giorgio (Southern Switzerland). In: Arratia G, Tintori A eds. *Mesozoic Fishes 3–Systematics, Paleoenvironments and Biodiversity* 3. 239–251
- Cavin L, 2010. Diversity of Mesozoic semionotiform fishes and the origin of gars (Lepisosteidae). *Naturwissenschaften*, 97: 1035–1040
- Ebert M, 2018. *Cerinichthys koelblae*, gen. et sp. nov., from the Upper Jurassic of Cerin, France, and its phylogenetic setting, leading to a reassessment of the phylogenetic relationships of Halecomorphi (Actinopterygii). *J Vert Paleont*, 38: e1420071
- Folkvord A, Hunter J R, 1986. Size-specific vulnerability of northern anchovy, *Engraulis mordax*, larvae to predation by fishes. *Fish Bull*, 84: 859–869
- Friedman M, 2015. The early evolution of ray-finned fishes. *Paleontology*, 58: 213–228
- Gardiner B G, 1960. A revision of certain actinopterygian and coelacanth fishes, chiefly from the Lower Lias. *Bull Br Mus Nat Hist Geol, Geology*, 4(7): 241–384
- Geng B H, Zhu M, Jin F, 2009. A revision and phylogenetic analysis of *Guizhoucoelacanthus* (Sarcopterygii, Actinistia) from the Triassic of China. *Vert Palasiat*, 47: 165–177
- Gill E L, 1923. The Permian fishes of the genus *Acentrophorus*. *Proc Zool Soc Lond*, 93: 19–40
- Grande L, 2010. An empirical synthetic pattern study of gars (Lepisosteiformes) and closely related species, based mostly on skeletal anatomy. The resurrection of holostei. *Copeia*, 10(Suppl): 1–871
- Grande L, Bemis W E, 1998. A comprehensive phylogenetic study of amiid fishes (Amiidae) based on comparative skeletal anatomy: an empirical search for interconnected patterns of natural history. *Soc Vert Paleont Mem (Suppl J Vert Paleont)*, 4: 1–690
- Herzog A, 2003. Eine Neubeschreibung der Gattung *Eoeugnathus*, Brough, 1939 (Actinopterygii; Halecomorphi) aus der alpinen Mitteltrias Graubündens. *Palaönt Zeit*, 77(1): 223–240
- Hurley I A, Mueller R L, Dunn K A et al., 2007. A new time-scale for ray-finned fish evolution. *Proc R Soc B*, 274: 489–498
- Jiang D Y, Motani R, Tintori A et al., 2020. Evidence supporting predation of 4-m marine reptile by Triassic megapredator.

iScience, 23: 101347

Jin F, 2001. Notes on the discovery of *Birgeria* in China. Vert PalAsiat, 39: 168–176

Karpouzi V S, Stergiou K I, 2003. The relationships between mouth size and shape and body length for 18 species of marine fishes and their trophic implications. J Fish Biol, 62: 1353–1365

Li C, Wu X C, Zhao L J et al., 2016. A new armored archosauriform (Diapsida: Archosauromorpha) from the marine Middle Triassic of China, with implications for the diverse life styles of archosauriforms prior to the diversification of Archosauria. Sci Nat, 103: 95

Li J L, 2006. A brief summary of the Triassic marine reptiles of China. Vert PalAsiat, 44: 99–108

Li Z G, Sun Z Y, Jiang D Y et al., 2016. LA-ICP-MS Zircon U-Pb age of the fossil layer of Triassic Xingyi Fauna from Xingyi, Guizhou, and its significance. Geol Rev, 62: 779–790

Liu G B, Yin G Z, Wang X H, 2002. On the most primitive amiid fish from Upper Triassic of Xingyi, Guizhou. Acta Palaeont Sin, 41: 461–463

Liu G B, Yin G Z, Wang X H et al., 2003. New discovered fishes from *Keichousaurus* bearing horizon of Late Triassic in Xingyi of Guizhou. Acta Palaeont Sin, 42: 346–366

Lombardo C, 2001. Actinopterygians from the Middle Triassic of northern Italy and Canton Ticino (Switzerland): anatomical descriptions and nomenclatural problems. Riv Ital Paleont Stratigr, 107: 345–369

López-Arbarello A, Ebert M, 2023. Taxonomic status of the caturid genera (Halecomorpha, Caturidae) and their Late Jurassic species. R Soc Open Sci, 10: 221318

López-Arbarello A, Sferco E, 2018. Neopterygian phylogeny: the merger assay. R Soc Open Sci, 5: 172337

López-Arbarello A, Sun Z Y, Sferco E et al., 2011. New species of *Sangiorgioichthys* Tintori and Lombardo, 2007 (Neopterygii, Semionotiformes) from the Anisian of Luoping (Yunnan Province, South China). Zootaxa, 2749: 25–39

Ma X Y, Xu G H, 2017. A new ionoscopiform fish (Holostei: Halecomorpha, Ionoscopiformes) from the Middle Triassic (Anisian) of Yunnan, China. Vert PalAsiat, 55: 92–106

Magnhagen C, Heibo E, 2001. Gape size allometry in pike reflects variation between lakes in prey availability and relative body depth. Funct Ecol, 15: 754–762

Metcalfe I, 2011. Palaeozoic–Mesozoic history of SE Asia. In: Hall R, Cottam M A, Wilson M E J eds. The SE Asian Gateway: History and Tectonics of the Australia–Asia Collision. Geol Soc, Spec Publ, London, 355: 7–35

Near T J, Eytan R I, Dornburg A et al., 2012. Resolution of ray-finned fish phylogeny and timing of diversification. Proc Nat Acad Sci, 109: 13698–13703

Olsen P E, 1984. The skull and pectoral girdle of the parasemionotid fish *Watsonulus eugnathoides* from the Early Triassic Sakamena group of Madagascar, with comments on the relationships of the holostean fishes. J Vert Paleont, 4: 481–499

Patterson C, 1973. Interrelationships of holosteans. In: Greenwood P H, Miles R S, Patterson C eds. Interrelationships of Fishes. Zool J Linn Soc, 53(Suppl): 233–305

Scharf S F, Juanes F, Rountree R A, 2000. Predator size–prey size relationships of marine fish predators: interspecific variation and effects of ontogeny and body size on trophic-niche breadth. Mar Ecol Prog Ser, 208: 229–248

Shang Q H, Wu X C, Li C, 2020. A new Ladinian nothosauroid (Sauropterygia) from Fuyuan, Yunnan Province, China. J Vert Paleont, 40: e1789651

- Su D Z, 1959. Triassic fishes from Kueichow, Southwest China. *Vert PalAsiat*, 3: 205–210
- Sun Z Y, Tintori A, Xu Y Z et al., 2016. A new non-parasemionotiform order of the Halecomorphi (Neopterygii, Actinopterygii) from the Middle Triassic of Tethys. *J Syst Palaeont*, 15: 223–240
- Swofford D L, 2003. PAUP*. Phylogenetic Analysis Using Parsimony (*and Other Methods). Version 4.0b10. Sunderland, Massachusetts: Sinauer Associates
- Tintori A, Lombardo C, 2007. A new early Semionotidae (Semionotiformes, Actinopterygii) from the Upper Ladinian of the Monte San Giorgio area (southern Switzerland and northern Italy). *Riv Ital Paleont Stratigr*, 113: 369–381
- Tintori A, Hitij T, Jiang D Y et al., 2014. Triassic actinopterygian fishes: the recovery after the end-Permian crisis. *Integr Zool*, 9: 349–411
- Tintori A, Sun Z Y, Lombardo C et al., 2015. Oldest stem Teleostei from the late Ladinian (Middle Triassic) of southern China. *Riv Ital Paleont Stratigr*, 121: 285–296
- Westoll T S, 1944. The Haplolepididae, a new family of Late Carboniferous bony fishes – a study in taxonomy and evolution. *Bull Am Mus Nat Hist*, 83: 1–121
- Xu G H, 2019. Osteology and phylogeny of *Robustichthys luopingensis*, the largest holostean fish in the Middle Triassic. *PeerJ*, 7: e7184
- Xu G H, 2020. A new species of Luganoia (Luganoiidae, Neopterygii) from the Middle Triassic Xingyi Biota, Guizhou, China. *Vert PalAsiat*, 58: 267–282
- Xu G H, Ma X Y, 2018. Redescription and phylogenetic reassessment of *Asialepidodus shiniensis* (Holostei: Halecomorphi) from the Middle Triassic (Ladinian) of China. *Zool J Linn Soc*, 184(1): 95–114
- Xu G H, Shen C C, 2015. *Panxianichthys imparilis* gen. et sp. nov., a new ionoscopiform (Halecomorphi) from the Middle Triassic of Guizhou, China. *Vert PalAsiat*, 53: 1–15
- Xu G H, Wu F X, 2012. A deep-bodied ginglymodian fish from the Middle Triassic of eastern Yunnan Province, China, and the phylogeny of lower neopterygians. *Chinese Sci Bull*, 57: 111–118
- Xu G H, Zhao L J, Gao K Q et al., 2012. A new stem-neopterygian fish from the Middle Triassic of China shows the earliest over-water gliding strategy of the vertebrates. *Proc R Soc B*, 280: 20122261
- Xu G H, Zhao L J, Coates M I, 2014. The oldest ionoscopiform from China sheds new light on the early evolution of halecomorph fishes. *Biol Lett*, 10: 20140204
- Xu G H, Gao K Q, Coates M I, 2015. Taxonomic revision of *Plesiofuro mingshuica* from the Lower Triassic of northern Gansu, China, and the relationships of early neopterygian clades. *J Vert Paleont*, 35(6): e1001515
- Xu G H, Ma X Y, Ren Y, 2018. *Fuyuanichthys wangi* gen. et sp. nov. from the Middle Triassic (Ladinian) of China highlights the early diversification of ginglymodian fishes. *PeerJ*, 6: e6054
- Young C C, 1958. On the new pachypleurosauroidea from Keichow, South-West China. *Vert PalAsiat*, 2: 69–81
- Zou X D, Balini M, Jiang D Y et al., 2015. Ammonoids from the Zhuganpo Member of the Falang Formation at Nimaigu and their relevance for dating the Xingyi Fossil-Lagerstaette (late Ladinian, Guizhou, China). *Riv Ital Paleont Stratigr*, 121: 135–161